ANNEX J TO SMALLPOX RESPONSE PLAN RESOURCES.

29 September 2002

APPENDIX J-1

Acronyms and Abbreviations.

ACAM - Acambis

ACC - Acute Care Center

ACIP - Advisory Committee on Immunization Practices

AE - Adverse Event

AFCITA - Air Force Complete Immunization Tracking Application

AFMAN – Air Force Manual

AFMLO - Air Force Medical Logistics Office

AFMOA - Air Force Medical Operations Agency

AHA – American Hospital Association

AIA - American Institute of Architects

AIDS - Acquired Immune Deficiency Syndrome

AIIR – Airborne Infectious Isolation Room (or area, AIIA)

AMA - American Medical Association

AMSA - Army Medical Surveillance Activity

AO - Area of Operations

AOI – Area of Interest

AOR – Area of Responsibility

AP - Aventis Pasteur

APHIS - Anaimal and Plant Health Inspection Service

APIC - Association for Professionals in Infection Control & Epidemiology

AHA- American Hospital Association

ASHE - American Society for Healthcare Engineering

ATI – Air Transportable Isolator (a chamber to contain patients with infectious diseases)

ATOC – Air Transportation Operations Center

BAT - Biological Augmentation Team

BioMedAC-Biological Medical Advisory Committee

BMBL - Biosafety in Microbiological and Biomedical Laboratories

BSC - BIOLOGICAL safety Cabinets

BSL - Bio-Safety Level

BPRP - Bioterrorism Preparedness & Response Program, CDC

BW- Biological Weapon/ Warfare

C-type Facility - (C for confirmed) Mode of shelter and care for people diagnosed with smallpox

CBRN - Chemical Biological Radiologic Nuclear

CBRNE – Chemical Biological Radiologic Nuclear Explosive

CDC - Centers for Disease Control & Prevention

CENTCOM - Central Command

CFR - Code of Federal Regulations

CINC - Commander-In-Chief

CIL- Critical Information Lists

CIO - Center/Institute/Office

CISA - Clinical Immunization Safety Assessment centers of excellence

CHPPM - Center for Health Promotion & Preventive Medicine

CM – Consequence Management

CMV - Cytomegalovirus

CONUS - Contiguous United States

CPR – Cardiopulmonary resuscitation

CXR – Chest radiograph X-ray

DA - Department of Army

DD - Department of Defense

DEERS - Defense Eligibility Enrollment Reporting System

DEPMEDS - Deployable Medical Systems

DIC – Disseminated Intravascular Coagulation

DGR - Dangerous Goods Regulations

DHHS - Department of Health & Human Services

DLA – Defense Logistics Agency

DMDC - Defense Manpower Data Center

DMAT – Disaster Medical Assistance Teams

DNA – Deoxyribonucleic Acid

DNIF – Duties Not Including Flying

DOC - Distribution Operations Center, USAMMA

DOT - Department of Transportation

DOMS- Directorate of Military Support

DoD - Department of Defense

DRG - Dangerous Goods Regulations

DSN - Defense Switch Network

DSS – Distribution Standard System

DVC - DynPort Vaccine Company

DVRD - Division of Viral & Rickettsial Diseases, CDC

DWI-Disaster Welfare Information

DQ - Division of Quarantine, CDC

EHBS - Enhanced Hospital-Based Surveillance

EISO - Epidemic Intelligence Service Officer

EPA - Environmental Protection Agency

Epi-Team - Epidemiologic Response Team

EPO - Epidemiology Program Office, CDC

EPRB - Emergency Preparedness & Response Branch, CDC

ESF – Emergency Support Function

ER - Emergency Room

ESSENCE - Electronic Surveillance System for the Early Notification of Community Based Epidemics

EUCOM - European Command

FBI - Federal Bureau of Investigations

FDA - Food & Drug Administration

FEMA - Federal Emergency Management Agency

FMP – Family Member Prefix

FPCON – Force Protection Condition

FOSC- Federal On-Scene Coordinator

FRP - Federal Response Plan

GCFS – Granulocyte Colony Stimulating Factor

GFVPRI - Generalized Febrile Vesicular-Pustular Rash Illness

GTN – Global Transportation Network

GVPRI - Generalized Vesicular-Pustular Rash Illness

HAD - Hospital-Approved Disinfectant

HCP - Health Care Providers

HCW - Healthcare Worker

HEPA - High-Efficiency Particulate Air (filter)

HHS - Health & Human Services, Department of

HICPAC – Hospital Infection Control Practices Advisory Committee

HIP - Hospital Infections Program, CDC

HIV/STD/TB - National Center for HIV, STD, & TB Prevention, CDC

HQ - Headquarters

HSRRB - Human Subjects Research Review Board (Army Surgeon General's IRB)

HSV - Herpes Simplex Virus

HVAC – Heating Ventilation and Air Conditioning

IATA - International Air Transportation Association

IAW - In Accordance With

ICAO – International Civil Aviation Organization

ICD9 – International Classification of Diseases, 9th edition

ICP - Infection Control Professionals

ICU - Intensive Care Unit

ICRA – Infection Control Risk assessments

ID - Infectious disease

IDSA - Infectious Diseases Society of America

IERA - Institute for ESOH (Environmental, Safety & Occupational Health) Risk Analysis,

USAF

ICU – Intensive Care Unit

IGIV - Immune globulin intravenous

IM - Intramuscular

IND - Investigational New Drug

IRB – Institutional Review Board (medical research ethics committee)

IV – Intravenous

IVIG - see IGIV

JAMA - Journal of the American Medical Association

JIC - Joint Information Center

JPMPG - Joint Preventive Medicine Policy Group

JS - Joint Staff

JSSED – Joint Service Sensitive Equipment Decontamination

JVAP - Joint Vaccine Acquisition Program, DoD

LACUC - Laboratory Animal Care & Use Committee

LRMC – Landstuhl Regional Medical Center

LFA – Lead Federal Agency

LRN - Laboratory Response Network

MADCP - Mortuary Affairs Decontamination Collection Point

MEDCEN - Medical Center (US Army)

MEDCOM - Medical Command (US Army)

MHS - Military Health System

MMWR - Morbidity & Mortality Weekly Report

MEDPROS - Medical Protection System (US Army software application)

MRO – Materiel Release Order

MTF - Military Treatment Facility

NACI - National Advisory Committee on Immunization, Canada

NAME – National Association of Medical Examiners

NAVMED - Naval Medical Command

NAVMEDLOGCOM - Navy Medical Logistics Command

NBC - Nuclear, Biological & Chemical

NCEH - National Center for Environmental Health, CDC

NCID - National Center for Infectious Diseases, CDC

NCR – National Capital Region

NDC - National Drug Code

NDMS- National Disaster Medical System

NEHC - Navy Environmental Health Center

NEO- Non Combatant Evacuation Operations

NEPMU-6 - Navy Environmental & Preventive Medicine Unit-6

NHRC - Naval Health Research Command, San Diego

NIOSH - National Institute for Occupational Safety & Health

NIP - National Immunization Program, CDC

NPS - National Pharmaceutical Stockpile, CDC

NPSB - National Pharmaceutical Stockpile Branch, CDC

NSN - National Stock Number

NYCBOH - New York City Board of Health

OASD(HA) - Office of the Assistant Secretary of Defense for Health Affairs

OASD(PA) - Office of the Assistant Secretary of Defense for Public Affairs

OCONUS - Outside the Contiguous United States

OEP - Office of Emergency Preparedness, DHHS

OHS - Office of Health & Safety, CDC

OGC - Office of General Council (either CDC or DoD)

OPCON – Under Operational Control

OSHA – Occupational Safety and Health Administration

OTSG - Office of the Surgeon General, US Army

PACOM - Pacific Command

PAG – Public Affairs Guidance

PAHO- Pan American Health Organization

PAO – Public Affairs Officers

PAPR – Powered Air Purifying Respirations

PAR - Population at Risk

PCR- Polymerase Chain Reaction(Laboratory test method)

PDF - Portable document format

PFU - Pock-forming units

PHA- Public Health Advisor

PHPPO- Public Health Practice Program Officer, CDC

PHS- Public Health Service

PI- Principal Investigator/Product Insert(Package insert)

PM- Preventive Medicine

PPE – Personal Protective Equipment\

ppm – parts per million

PV- Prime Vendor

PAHO - Pan-American Health Organization

PAO - Public Affairs Office or Officers

PCR – Polymerase Chain Reaction (laboratory test method)

PHA - Public Health Advisor

PHPPO - Public Health Practice Program Office, CDC

PHS - Public Health Service

PI - Principal investigator, product insert (package insert)

PM - Preventive Medicine

PPE – Personal Protective Equipment

PV – Prime Vendor

R-type Facility - Residential mode of housing for surveillance of vaccinated contacts of smallpox cases

RC - Reserve Component

ROM – Restriction of Movement

RUC - Reporting Unit Codes

RMW - Regulated Medical Waste

RRAT - Rapid Response & Advanced Technology Laboratory, CDC

SACEUR- Supreme Allied Commander- Europe

SAMS - Shipboard Automated Medical System

SHEA – Society of Healthcare Epidemiologists of America

SIR - Serious Incident Report

SMART - Special Medical Augmentation Response Team, US Army

SOFA - Status Of Forces Agreement

SRT - Smallpox Response Teams

TAML - Theater Army Medical Laboratory

TEU - Technical Escort Unit

T-TEAMS – Treatment Teams

TYCOM – Type Commander (Naval)

Type C Facility - (C for confirmed) Mode of shelter and care for people diagnosed with smallpox

Type R Facility - Residential mode of housing for surveillance of vaccinated contacts of smallpox cases

Type X Facility - (X for Uncertain) Mode of shelter for surveillance of contacts of smallpox cases with fever but without signs and symptoms diagnostic of smallpox

UI - Unit of Issue

UIC - Unit Identification Code

UPS - United Parcel Service

URL - Universal Resource Locator

USAMMA - United States Army Medical Materiel Agency

USAMMDA - United States Army Medical Materiel Development Activity

USACHPPM - United States Army Center for Health Promotion & Preventive Medicine

USAMRIID - United States Army Medical Research Institute of Infectious Diseases

USAMRMC – United States Army Medical Research and Materiel Command

USC - United States Code

USDA – United States Department of Agriculture

USPS - United States Postal Service

USUHS - Uniformed Services University of the Health Sciences

UV - Ultraviolet

VAERS - Vaccine Adverse Event Reporting System

VEHB - Viral Exanthems & Herpesvirus Branch, CDC

VIG - Vaccinia Immune Globulin

VHC - Vaccine Healthcare Center

WMD - Weapons of Mass Destruction

WHO - World Health Organization

WRAMC - Walter Reed Army Medical Center

X-type Facility - (X for Uncertain) Mode of shelter for surveillance of contacts of smallpox cases with fever but without signs and symptoms diagnostic of smallpox

APPENDIX J-2 URLs for CDC Smallpox Response Plan Document. http://www.bt.cdc.gov/DocumentsApp/Smallpox/RPG/index.asp

	Source of Exposure		
Docu- ment	Title / Subject	PDF URL	HTML URL
	Executive Summary plus parts II through VI	http://www.bt.cdc.gov/DocumentsApp/Sma llpox/RPG/FrontPgs/Sections-i-iv.pdf	http://www.bt.cdc.gov/DocumentsApp/Sma llpox/RPG/FrontPgs/Sections-i-iv.doc
CDC Guide A	Surveillance, Contact Tracing, and Epidemiological Investigation	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a.pdf	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a.doc
	Smallpox Case Investigation		
Form 1A	Page 1 of 2	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a-form-1a-pg1.pdf	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a-form-1a-pg1.ppt
Form 1A	Page 2 of 2	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a-form-1a-pg2.pdf	http://www.bt.cdc.gov/DocumentsApp/Sma llpox/RPG/guideA/guide-a-form-1a-pg2.ppt
Form 1B	Page 1 of 2	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a-form-1b-pg1.pdf	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a-form-1b-pg1.ppt
Form 1B	Page 2 of 2	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a-form-1b-pg2.pdf	http://www.bt.cdc.gov/DocumentsApp/Sma llpox/RPG/guideA/guide-a-form-1b-pg2.ppt
	Contact Tracing		
Form 2	Interviewer Checklist / Contact Information	http://www.bt.cdc.gov/DocumentsApp/Sma llpox/RPG/guideA/guide-a-form-2.pdf	http://www.bt.cdc.gov/DocumentsApp/Sma llpox/RPG/guideA/guide-a-form-2.doc
Form 2a	Case Travel / Activity Calendar	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a-form-2a.pdf	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a-form-2a.doc
Form 2b	Interviewer Contact / Site Summary Worksheet	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a-form-2b.pdf	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a-form-2c.doc
Form 2c	Contact Transportation	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a-form-2c.pdf	http://www.bt.cdc.gov/DocumentsApp/Sma Ilpox/RPG/guideA/guide-a-form-2c.doc
Form 2d	Out of Area Travel Log	http://www.bt.cdc.gov/DocumentsApp/Sma llpox/RPG/guideA/guide-a-form-2d.pdf	http://www.bt.cdc.gov/DocumentsApp/Sma llpox/RPG/guideA/guide-a-form-2d.xls

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Form 4	Source of Exposure	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
		allpox/RPG/guideA/guide-a-form-4.pdf	Ilpox/RPG/guideA/guide-a-form-4-now-3.doc
Form 6	Daily Case Status	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
1 01111 0	Tracking	allpox/RPG/guideA/guide-a-form-6.pdf	Ilpox/RPG/guideA/guide-a-form-6.xls
Form 7	Daily Case Status	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
	Tracking	allpox/RPG/guideA/guide-a-form-7.pdf	Ilpox/RPG/guideA/guide-a-form-7.doc
Form 8	Contact Interview	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
		allpox/RPG/guideA/guide-a-form-8.pdf	Ilpox/RPG/guideA/guide-a-form-8.doc
Form 9	Contact Vaccination	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
	Referral	allpox/RPG/guideA/guide-a-form-9.pdf	<u>Ilpox/RPG/guideA/guide-a-form-9.doc</u>
Form 10	Individual Contact	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
	Surveillance	allpox/RPG/guideA/guide-a-form-10.pdf	<u>Ilpox/RPG/guideA/guide-a-form-10.doc</u>
Form 11	Contact Tracking (Daily	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
	Record Master form)	allpox/RPG/guideA/guide-a-form-11.pdf	<u>Ilpox/RPG/guideA/guide-a-form-11.doc</u>
CDC	Vaccination Guidelines		
Guide B	for State and Local		
	Health Agencies		
	Part 1 (pages 1 to 12)	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
		allpox/RPG/GuideB/guide-b-part1of2.pdf	<u>Ilpox/RPG/GuideB/guide-b-part1of2.doc</u>
	Part 2 (pages 13 to 19)	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
		allpox/RPG/GuideB/guide-b-part2of2.pdf	<u>Ilpox/RPG/GuideB/guide-b-part2of2.doc</u>
CDC Guide C	Isolation and Quarantine Guidelines		
	Part 1 (pages 1 to 18)	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
	, ,	allpox/RPG/GuideC/guide-C-pages1-	Ilpox/RPG/GuideC/guide-C-pages1-
		18only.pdf	18only.doc
	Part 2 (pages 1 to 21)	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
	, ,	allpox/RPG/GuideC/guide-C-pages19-	Ilpox/RPG/GuideC/guide-C-pages19-
		21only.pdf	21only.doc
CDC	Specimen Collection	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
Guide D	and Transport	allpox/RPG/GuideD/Guide-D.pdf	Ilpox/RPG/GuideD/Guide-D.doc
	Guidelines		

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CDC	Communications Plans	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
Guide E	and Activities	allpox/RPG/GuideE/Guide-E.pdf	Ilpox/RPG/GuideE/Guide-E.doc
CDC	Decontamination	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
Guide F	Guidelines	allpox/RPG/GuideF/Guide-F.pdf	Ilpox/RPG/GuideF/Guide-F.doc
CDC	Annex 1: Overview of		
Annex 1	Smallpox, Clin. Present-		
	ation, Medical Care of		
	Smallpox Patients		
	Part 1 (pages 1 to 9)	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
		allpox/RPG/annex/annex-1-part1of3.pdf	<u>Ilpox/RPG/annex/annex-1-part1of3.doc</u>
	Part 2 (pages 10 to 16)	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
		allpox/RPG/annex/annex-1-part2of3.pdf	<u>Ilpox/RPG/annex/annex-1-part2of3.doc</u>
	Part 3 (pages 17 to 19)	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
		allpox/RPG/annex/annex-1-part3of3.pdf	<u>Ilpox/RPG/annex/annex-1-part3of3.doc</u>
CDC	Guidelines for Smallpox	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
Annex 2	Vaccination Clinics	allpox/RPG/annex/annex-2.pdf	<u>Ilpox/RPG/annex/annex-2.doc</u>
CDC	Vaccine Adverse Event	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
Annex 3	Reporting	allpox/RPG/annex/annex-3.pdf	Ilpox/RPG/annex/annex-3.doc
CDC	Suggested Pre-Event	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
Annex 4	Activities for State &	allpox/RPG/annex/annex-4.pdf	Ilpox/RPG/annex/annex-4.doc
	Local Health Authorities		
	Generalized Vesicular or	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
	Pustular Rash Illness	allpox/RPG/annex/annex-4-rash-color.pdf	Ilpox/RPG/annex/annex-4-rash-color.ppt
	Protocol		
CDC	Glossary of	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
Annex 5	Abbreviations and	allpox/RPG/annex/annex-5.pdf	Ilpox/RPG/annex/annex-5.doc
	Smallpox References		
CDC	Miscellaneous Forms in	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
Annex 6	development	allpox/RPG/annex/annex-6.pdf	Ilpox/RPG/annex/annex-6.doc
CDC	Checklists for State/	http://www.bt.cdc.gov/DocumentsApp/Sm	http://www.bt.cdc.gov/DocumentsApp/Sma
Annex 7	Local/ CDC Personnel	allpox/RPG/annex/annex-7.pdf	<u>Ilpox/RPG/annex/annex-7.doc</u>
	Actions in a Smallpox		
	Emergency		

APPENDIX J-3

Professional Resources on Smallpox.

Agency for Healthcare Research and Quality Bioterrorism Education Site. www.bioterrorism.uab.edu/

American College of Physicians and American Society of Internal Medicine (ACCP-ASIM), Bioterrorism Resource Center, www.acponline.org/bioterro/index.html

American Society for Microbiology (ASM), Resources Related to Biological Weapons Control and Bioterrorism Preparedness, www.asmusa.org/pcsrc/bioprep.htm

Association for Infection Control and Epidemiology, Inc. (APIC), (202) 789-1890, (202) 789-1890, http://www.apic.org/bioterror/, www.apicelearn.org

Centers for Disease Control and Prevention, http://www.bt.cdc.gov/ or http://www.cdc.gov/, (404) 639-3311

Centers for Disease Control and Prevention (CDC). Facts about smallpox., www.bt.cdc.gov/DocumentsApp/FactSheet/SmallPox/About.asp

Centers for Disease Control and Prevention (CDC). Frequently asked questions, www.bt.cdc.gov/DocumentsApp/SmallPox/10242001fags/10242001SmallpoxFAQs.asp

CDC Responds series: Smallpox: What Every Clinician Should Know. www.phppo.cdc.gov/phtn/default.asp

CDC Public Health Emergency & Response site, <u>www.bt.cdc.gov</u>, cdc.gov/ncidod/diseases/bioterr.htm

Department of Defense, http://www.defenselink.mil/, (703) 697-5737

DoD; Improving Local And State Agency Response To Terrorist Incidents Involving Biological Weapons.

http://www2.sbccom.armv.mil/hld/downloads/bwirp/bwirp interim planning guide.pdf

Department of Health and Human Services, http://www.dhhs.gov/, 1-877-696-6775

Domestic Preparedness Helpline: 1-800-368-6498

Domestic Preparedness Website: http://www.nbc-prepare.org/

Environmental Protection Agency, http://www.epa.gov/, (202) 260-2090

Federal Bureau of Investigation, http://www.fbi.gov/, (202) 324-3000

Federal Emergency Management Agency, http://www.fema.gov/, (202) 646-4600

Infectious Disease Society of America (IDSA), Bioterrorism Preparedness (includes links to useful articles), www.idsociety.org/PA/PS&P/BT Preparedness 10-2-01.htm

Johns Hopkins Center for Civilian Biodefense Studies, http://www.hopkins-biodefense.org/pages/agents/agentsmallpox.html

National Domestic Preparedness Office, http://www.ndpo.gov/, (202) 324-9026

Society for Healthcare Epidemiology of America, Inc., http://www.shea-online.org/BTprep.html

University of Alabama – Birmingham, Emerging Infections and Potential Bioterrorist Agents, www.bioterrorism.uab.edu/

UCLA Department of Epidemiology, www.ph.ucla.edu/epi/bioter/bioterrorism.html

U.S. Army Medical Research Institute of Infectious Diseases (USAMRIID), www.usamriid.army.mil

USAMRIID's Medical Management of Biological Casualties Handbook, www.usamriid.army.mil/education/bluebook.html

US Army Medical NBC Information Server, www.nbc-med.org/others/Default.html

US Army NBC Information Server, www.nbc-med.org/others/Default.html

APPENDIX J-4

Reference Publications on Smallpox.

Advisory Committee on Immunization Practices. Vaccinia (smallpox) vaccine. *MMWR* 2001;50(RR-10):1-25. http://www.cdc.gov/mmwr/PDF/rr/rr5010.pdf. See Appendix J-7.

Advisory Committee on Immunization Practices. Draft supplemental recommendations on smallpox (vaccinia) vaccine. 2002 June 20. www.cdc.gov/nip/smallpox/supp recs.htm

Atkinson W, Wolfe C, Humiston S, Nelson R, ed. Chapter 18, Smallpox. In: *Epidemiology & Prevention of Vaccine-Preventable Diseases*, 7th ed. Atlanta: Centers for Disease Control & Prevention, Apr 2002. http://www.cdc.gov/nip/publications/pink/#download.

Barbera J, Macintyre A, Gostin L, Inglesby T, O'Toole T, DeAtley C, Tonat K, Layton M. Large-scale quarantine following biological terrorism in the United States: Scientific examination, logistic and legal limits, and possible consequences. *JAMA* 2001;286:2711-2717. http://jama.ama-assn.org/issues/v286n21/rpdf/jsc10254.pdf

Breman JG, Henderson DA. Diagnosis and management of smallpox. *N Engl J Med* 2002; 346:1300-1308. http://content.nejm.org/cgi/reprint/346/17/1300.pdf.

Canadian National Advisory Committee on Immunisation. Statement on smallpox vaccination. *Can Comm Dis Rep* 2002;28(ACS-1):1-12. http://www.hc-sc.gc.ca/pphb-dgspsp/publicat/ccdr-rmtc/02pdf/acs28-1.pdf.

Coates JB Jr., Hoff EC. Medical Department, United States Army, in World War II— Preventive Medicine in World War II, Volume III: Personal Health Measures and Immunization. Washington, DC: Department of the Army, 1955, pages 280-287, 351-352.

Coates JB Jr., Hoff EC. Medical Department, United States Army, in World War II— Preventive Medicine in World War II, Volume VII: Communicable Diseases— Arthropodborne Diseases Other Than Malaria. Washington, DC: Department of the Army, 1986, pages 17-18.

Fenner F, Henderson DA, Arita I, Jezek Z, Ladnyi ID. *Smallpox and Its Eradication*. Geneva: World Health Organization, 1988. http://www.who.int/emc/diseases/smallpox/Smallpoxeradication.html.

Frey SE, Newman FK, Cruz J, Shelton WB, Tennant JM, Polach T, Rothman AL, Kennedy JS, Wolff M, Belshe RB, Ennis FA. Dose-related effects of smallpox vaccine. *N Engl J Med* 2002;346:1275-1280. http://content.nejm.org/cgi/reprint/346/17/1275.pdf.

Frey SE, Couch RB, Tacket CO, Treanor JJ, Wolff M, Newman FK, Atmar RL, Edelman R, Nolan CM, Belshe RB, National Institute of Allergy and Infectious Diseases Smallpox Vaccine Study Group. Clinical responses to undiluted and diluted smallpox vaccine. *N Engl J Med* 2002;346:1265-1274. http://content.nejm.org/cgi/reprint/346/17/1265.pdf.

Gani R, Leach S. Transmission potential of smallpox in contemporary situations. *Nature* 2001;414:748-751; errata 415:1056.

Haim M, Gdalevich M, Mimouni D, Askenazi I, Shemer J. Adverse reactions to smallpox vaccine: The Israel Defense Force experience, 1991 to 1996: A comparison with previous surveys. *Milit Med* 2000;165:287-289.

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APPENDIX J-5

Restriction of Movement as a Tool for the Control of Contagious Disease on the Battlefield.

Adapted from: Joint Venture Oversight Group (a joint US/UK bilateral collaboration), November 2001.

Introduction

- 1. Military personnel face a serious threat from disease during military operations. Historically, disease, not warfare, was the greatest source of casualties on the battlefield. Unchecked, casualties from disease have the potential to seriously degrade the operational effectiveness of deployed forces or, in the worst case, endanger the achievement of mission goals.
- 2. Disease in humans originates from some naturally occurring micro-organisms such as bacteria and viruses. Some of these organisms are highly contagious (i.e., have the capacity to spread from person to person). Others are infrequent causes of illness in man in normal conditions (e.g., anthrax), but may be highly contagious when introduced to the environment deliberately (e.g., in a biological weapon, BW). The control of contagious diseases presents particular problems in a military operational setting. Warfare inevitably involves large numbers of personnel working and living closely together for extended periods of time under stressful conditions. In these circumstances, a contagious disease has the potential to spread rapidly whether due to poor public or environmental health and hygiene or as a result of introduction by release from a biological weapon.

Principles of Disease Control

- 3. Whatever its source, contagious disease on the battlefield must be managed and controlled in order to maintain the operational effectiveness of forces. Modern medical science, public-health practices and good environmental health management provide many effective disease control tools.
- 4. Preventing, or controlling, the spread of contagious disease is accomplished by rendering those at risk resistant to the disease and limiting their exposure. In general, this is effected by medical intervention, such as immunization, or by restricting contact between healthy individuals and sources of the disease (other individuals, animals, insects etc.).
- 5. This guide focuses on one tool available to a Commander for the control of contagious disease Restriction of Movement (ROM). The use of ROM will present him with a number of unique challenges and dilemmas. However, in reading this guide, it is important to bear in mind that contagious disease control relies on the co-ordinated use of a number of techniques.

What is Restriction of Movement (ROM)?

- 6. ROM is a tool for maintaining operational effectiveness in the face of a contagious disease, whether natural or artificial (for example, a biological-weapon attack). It aims to control the spread of the disease by restricting contact between healthy groups of personnel and those who have, or are suspected of having, contracted it. Personnel covered by ROM do not necessarily need to be removed from operations; wherever possible it should be implemented in such a way as to allow them to continue their mission. ROM may also be necessary to reduce the risk that a contagious disease is transferred back to the home base. A separate paper ("Decision making tool for the evacuation of military casualties following suspect BW exposure") provides guidance on the evacuation of casualties following a suspected or confirmed attack using BW.
- 7. A note on terminology: The terms 'quarantine' and 'isolation' are often used in the context of preventing contact between healthy populations and those either infected, or suspected of being infected, with a contagious disease. Quarantine involves the detention of an individual, or group, who is suspected of having been exposed to a contagious disease, until it is deemed that they have escaped infection. Isolation is the separation of an infected individual from a healthy population. Both rely on restricting the movement of individuals to some degree. During military operations where personnel have contracted, or are suspected of having been exposed to, a contagious disease, a Commander may need to consider using either, or both. Throughout this guide we will therefore use the more universal term, restriction of movement, or ROM.

Overview of Approach to Disease Control

- 8. Before considering ROM in detail, it is necessary to understand the overall approach to disease control in a theatre of operations (see figure). Before deployment, general Nuclear, Biological & Chemical (NBC) and medical **intelligence** and **reconnaissance** is required in order to assess threats to health of the deploying force from disease. This information forms an integral part of the intelligence preparation of the battlespace and will be incorporated into the operational estimate. It provides a baseline against which to assess any subsequent disease outbreaks. This may be particularly important in assessing whether an outbreak is naturally occurring rather than the result of a biological attack. On the basis of the resulting **health risk assessment**, medical staff will advise the Commander on the **force health protection** options available to him. The aim should be to reduce the risk of disease exposure to as low as reasonably practical. Typical steps that the Commander will need to consider prior to, and during, deployment include:
- Appropriate theatre surveillance measures to ensure early detection of potential exposures to disease. This includes pre-deployment surveillance.
- Measures to avoid, or minimize, exposure to potential sources of disease. In many cases, operational considerations will make this difficult, in which case the steps

below should be taken in order to minimize the risks.

- Ensuring that all personnel receive appropriate pre-deployment health advice and training concerning the in-theatre disease risks and the measures required to minimize the risks of exposure.
- Ensuring that personnel are offered appropriate medical countermeasures (for example, immunization) prior to deployment and that adequate supplies are available for use in-theatre as required.

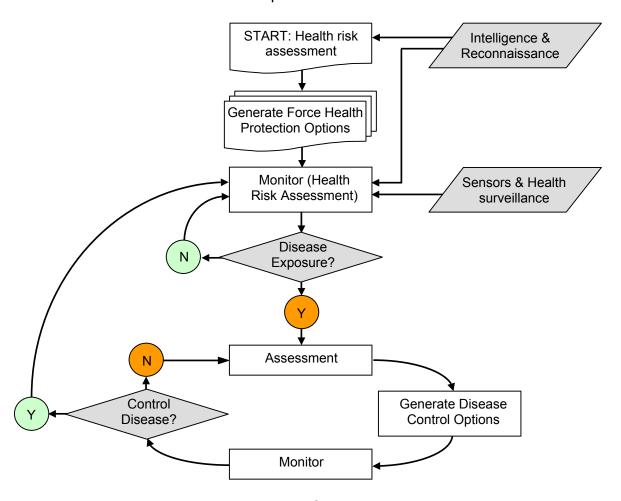


Figure: Overall approach to in-theatre disease control.

- 9. During deployment, the risks to the health of personnel are continuously **monitored** to ensure force health protection. Intelligence and reconnaissance continue to play an important role, and medical staff will provide the Commander with a health risk assessment. The first indication that personnel have been exposed to disease will be from one of two sources:
- Health surveillance and morbidity analysis picking up unusually high numbers, or distributions of illness amongst personnel, local civilians or others in theatre (e.g.

NGOs).

- Biological Warfare (BW) sensor alarms.
- 10. It may take some time to establish whether personnel have been exposed to a disease-causing agent, what that agent is and whether it is contagious. <u>Until a definitive diagnosis or identification can be made, the Commander should assume that personnel have been exposed to, and infected by, a contagious disease and take all possible steps, balanced against wider operational imperatives, to limit the spread of that <u>disease</u>. It may, in the initial stages, also be difficult to distinguish between a naturally occurring disease event and one as the result of a BW attack. All disease events will therefore require careful investigation involving a variety of medical and NBC specialists.</u>
- 11. Based on the evidence available and medical advice, the Commander must, as a matter of priority, **assess** the likely extent of exposure and infection and then in close consultation with the medical staff, taking the operational situation into account, generate **options** for controlling the disease event. These might include:
- Removing individuals showing disease symptoms for treatment as soon as possible.
- Instructing personnel within affected areas (or zones) on steps to minimize disease spread. Depending on the circumstances these might include decontamination procedures and environmental health controls such as enhanced monitoring of food and water supplies, more stringent hygiene measures, removing personnel from possible disease sources etc.
- Distributing, and instructing personnel to take, available medical countermeasures.
- Enhancing health surveillance of personnel within the suspected areas of exposure.
- Taking steps to minimize contact between personnel in the affected and nonaffected areas by the use of ROM (see below).
- 12. A period of continuous **monitoring** and **assessment** follows during which the success of **disease control measures** are monitored and progressively refined or modified. A high priority must be to obtain a definitive diagnosis or identification of the disease-causing agent so that the Commander and medical staff are able to tailor control measures to that particular pathogen and its mode of transmission. In particular, ROM will not be necessary for diseases that are not contagious, and may not be necessary for diseases against which physical protection (such as oro-nasal masks), or available medical countermeasures, provide an adequate level of protection.

Implementing Restriction of Movement (ROM)

13. A Commander will need to consider ROM following evidence that personnel have been exposed to disease. As described at Appendix J-5, paragraph 9, this evidence could come from BW sensors or health surveillance. Plans of action for these two eventualities are given at addendum A and addendum B respectively. Further considerations in implementing ROM are below.

General Operational Considerations

- 14. If the use of ROM is contemplated at any stage during an operation, it must be with the full knowledge that the impact on operational effectiveness is likely to be significant. The only stage of an operation when ROM is unlikely to play a significant deleterious role is during the close of an operation when personnel are being returned home. Here, ROM would be aimed not at preserving the fighting integrity of the force but, rather, reducing the risk of introducing contagious disease into the home base.
- 15. The implementation of ROM will restrict the ability of the Commander to use affected force elements and the fighting effectiveness of those units. There is also the danger that a perception on the part of an aggressor that ROM is likely to be used in response to a BW attack may make the use (or threatened use) of biological agents more attractive. In practice, the operational impact of disease control measures will need to be balanced against the potential consequences of the spread of a contagious disease. Operational pressures may dictate a policy that accepts the limited spread of a contagious disease because the implementation of ROM would result in the loss of the military objective. Special attention will need to be paid to highly mobile battlefield assets such as logistical units and Special Forces since these could inadvertently spread a contagious disease widely across the force structure or back to the home base.

Scale of Restriction of Movement (ROM)

- 16. The scale of ROM will depend on the precise set of circumstances surrounding any decision to implement it. It is most likely that ROM will be based around a particular geographic area or group of personnel. Personnel who may have been exposed to a BW agent should be instructed to carry out standard BW decontamination procedures. Most BW agents decay quickly when exposed to sunlight and drying, so contamination of the environment is unlikely to pose a significant hazard for an extended period (days or weeks) following an attack. Definition of ROM by geographic area is therefore unlikely to be useful beyond defining those force elements that may have been exposed to an agent. The general guideline is that, wherever possible, ROM should cover autonomous force elements so that they are able to continue their operational tasks while minimizing their contact with other unaffected units. This preserves the integrity of the chain of command, which will be critical both for ensuring that ROM is properly enforced and communication (see below).
- 17. One priority, whether or not ROM is used, is to ensure that any contagious disease is not spread back to the home base. As a minimum, it will be necessary to ensure that those returning home from a theatre of operation where a contagious disease is confirmed, or suspected, are the subjects of rigorous follow-up health surveillance. Special procedures

may be need to be implemented for personnel and assets that cycle in and out of theatre with high frequencies, for example strategic lift aircrew.

Treatment of Civilians

18. Civilians play an increasing role in the support of military operations. It is therefore very likely that the use of ROM will also affect civilians. DoD civilian workers and contractors likely will be treated in the same fashion as military personnel. However, where ROM affects other civilians (e.g. those of a host nation) its imposition and enforcement will be particularly problematic – for example, it may be necessary to segregate forces from the local populace. Close liaison with host nation authorities will be necessary in order to implement ROM successfully.

Duration of ROM

19. Once the decision to implement ROM has been taken, the criteria under which it will be lifted should also be identified. These 'exit criteria' will need to be based on medical advice and will primarily depend on the biological agent involved. The duration of ROM might be tailored to the projected incubation period of the disease following infection. As a minimum, this period should elapse without cases of disease occurring amongst personnel before ROM can be lifted. However, caution is needed since projections about the incubation time of contagious diseases are often imprecise and based on observations of naturally occurring forms of the disease. They may also be modified when biological agents are used as BW weapons or when personnel have used medical countermeasures. Once ROM is lifted, additional health surveillance for personnel should continue for an extended period to ensure that any subsequent outbreak, or re-emergence, of disease is quickly detected.

Morale & Communication

- 20. The use of ROM is likely to have far reaching psychological consequences on the personnel directly affected, the local civilian population and the perceptions of those at home. The successful prosecution of the operation may depend on dealing with the very natural feelings of fear, confusion, panic, indignation and anger likely to be displayed by different groups.
- Personnel affected by ROM. The imposition of ROM is likely to have a severe psychological impact on those groups who are directly affected, particularly if some of their number begin to fall ill. As suggested above, ROM should, wherever possible, be based around autonomous force elements that can continue some, or all, of their operational activities. This will ensure that personnel are focussed on continuing to perform their operational tasks rather than the implications of ROM. Clear communication is key and this approach also preserves the chain of command which has a crucial role to play in this respect. A decision to withhold information in order to reduce anxiety, although well intentioned, is likely to be counterproductive since rumour and confusion will spread quickly. Clear, concise and realistic messages should therefore be conveyed by a skilled health risk communicator as

soon as possible. These should cover:

- Why ROM has been implemented;
- The threat or risk faced by personnel factual, realistic assessments of the risks faced by personnel (including any details of the agents that might be involved);
- What precisely ROM involves clear explanation of how it will affect personnel and their operational activities;
- Other measures being taken including the delivery of medical assistance (medicines and treatment).
- When ROM will be lifted.
- The Main Force. Knowledge that ROM has been implemented on a component of the force is likely to be damaging. Again, clear and timely information will be vital and should be along the same lines as that for those directly affected.
- Local Civilian Population. Knowledge that a contagious disease is a problem amongst a locally deployed force has the potential to cause panic and disorder. Extremely close liaison with host nation civil and military authorities will be required to ensure that the situation is contained. All communication with the local civilian population should employ risk-communication principles.
- The Home Base. The media, perhaps as the result of a BW attack, will seize on news that personnel have been exposed to disease. Outrage, frustration, demands for action (and retaliation in the case of a BW attack), perhaps even calls for withdrawal will quickly follow. Many will fear the disease being spread back home. Relatives will be particularly anxious for news and information that the movements of apparently healthy personnel are being deliberately restricted, perhaps placing them at greater risk, will not be well received. The news media will need careful and professional handling. Commanders should take advice on this as soon as a decision to implement ROM is taken. Key messages will need to cover:
- Why ROM is necessary a broad explanation of the threat/risks faced by personnel;
- The aim of ROM protecting personnel and ensuring the success of the operation;
- Ensuring that any casualties get the best possible treatment.
- Ending ROM.

Summary

- The basic principles of the control of contagious disease on the battlefield are broadly the same whether the disease is naturally occurring or artificially introduced.
- ROM is <u>one</u> tool that may be considered by the Commander during an operation in order to control the spread of an infection amongst personnel. It should be used in concert with other measures and its use needs to be carefully balanced against its operational impact.
- Where ROM is implemented, it should, as far as possible, involve autonomous units that preserve the chain of command. Clear, precise and timely communication to personnel affected is critical.

Useful References

Joint NBC Defence Doctrine JWP 3-61
Joint Medical Doctrine JWP 4-03
Joint Doctrine for Operations in NBC Environments JP 3-11

ADDENDUM A: Evidence from BW Sensor Alarms

In this case, the priority will be to establish whether a biological weapons attack has taken place but, until confirmed otherwise, the Commander should assume that personnel have been exposed to a contagious disease. Action should be taken as follows:

- STEP 1: Carry out standard BW defence procedures. Establish the zone(s) within which it is most likely that personnel have been exposed to the BW agent (or, alternatively, where exposure is unlikely to have taken place).
- STEP 2: With medical staff, immediately review force health protection options with the assumption that the disease is contagious. See Appendix J-5, paragraph 11.
- STEP 3: Identify the agent used. This may be accomplished either by direct detection by BW sensors or may require the analysis of samples from the environment, or personnel, in a laboratory.
- STEP 4: Once the agent is identified, seek medical advice and review and refine the disease control measures taken at STEP 2, including the use of ROM. For all contagious disease there is a delay between exposure and the point at which infected individuals become contagious (i.e. are capable of passing the disease to others). This period depends on the disease, but ranges from one to several days. During this period personnel who have been exposed may have dispersed to different locations and may be difficult to track. A Commander therefore has up to 24 hours following initial detection in order to seek advice, reach a decision and, if necessary, prepare for the introduction of ROM, but should aim to do so as soon as possible. This process should include the generation of criteria for lifting ROM (see Appendix J-5, paragraph 19). These exit criteria will be based on the typical incubation period following exposure for the disease caused by the agent. For example, it may be possible to consider lifting ROM if no cases occur within several days of initial detection.
- STEP 5. Implement ROM (if required) and other disease control measures. In the initial instance, ROM is likely to be based on the zones identified at STEP1, but needs to take into account operational practicalities. For example, where only part of a unit has been exposed to a contagious agent, it may make more sense to apply ROM to the whole unit. Units within this area can continue to operate, but contact with other elements of the force should be limited as far as possible. However, the operational situation will determine whether unexposed personnel should be deployed in a ROM area.
- STEP 6. Enhance health surveillance and monitor success of control measures. This continues until the exit criteria identified at STEP 3 are satisfied (in which case

ROM can be lifted) or the first cases of disease amongst personnel are identified (proceed to STEP 1 in Addendum B).

ADDENDUM B: Evidence from Health Surveillance

In this case, the fact that personnel have been infected by disease is likely to become apparent over a period of days, even weeks. Even where the introduction of disease takes place as the result of a single event (e.g. the use of a single biological weapon) as well as being spread over a long period of time, cases may have a large geographic distribution caused by the movement of personnel and secondary spread of infection. A decision to implement ROM will be more difficult in these circumstances because the seriousness and extent of any disease outbreak may not be clear. If health surveillance indicates that personnel may have been infected, action should be as follows:

STEP 1. Prompt isolation and treatment of those with the symptoms of disease. Consideration will need to be given to how medical support will be delivered to the sick. It may be feasible to remove casualties, with appropriate precautions, to field hospitals, but where large numbers of personnel are affected this may not be possible. In this case medical facilities may need to be deployed into ROM areas.

STEP 2. With medical staff, immediately review force health protection options. See Appendix J-5, paragraph 11.

STEP 3. Definitive identification of the agent responsible. This will be arrived at based on medical diagnosis of the symptoms of infected individuals and laboratory analysis of patient samples. This may require the evacuation of a single case, under rigorously controlled conditions, back to the UK and/or US.

STEP 4. Once the agent is identified, seek medical advice and review and refine the disease control measures taken at STEP 2, including the use of ROM. If the agent does not cause a contagious disease, ROM is not required. If, however, the agent is confirmed as contagious the introduction of ROM will need to be considered (if not already implemented in STEP 2) to limit any potential spread amongst healthy personnel. In this case, the imperative will be to make an early decision concerning whether or not to use ROM, since the disease is already spreading amongst personnel. In the case of a highly contagious agent, delay may make the situation far worse than it may otherwise have been. Criteria for the lifting of ROM will also need to be generated (see Appendix J-5, paragraph 19).

STEP 5. Implement ROM (if required) in concert with other control measures. Here ROM is likely to be based on individuals or groups who have come into close contact with confirmed cases. As the disease runs its course, the groups affected by ROM may need to be expanded, or reduced. The operational impact of ROM can be minimized by basing it around autonomous force elements that are able to sustain their military functions.

STEP 6. Enhance health surveillance and monitor success of control measures.

STEP 7. Based on exit criteria, lift ROM. This may take place progressively as units, or force elements are confirmed disease free.

APPENDIX J-6

Considerations In Air-Evacuation Of A Smallpox Patient from Overseas

Adapted from: Decision Making Tool For Evacuation Of Military Casualties Following Suspect BW Exposure, Joint Venture Oversight Group (a joint US/UK bilateral collaboration), November 2001.

- 1. Introduction. Patients with an infection caused by a biological warfare (BW) attack can be medically managed in the same manner as those with a disease caused by natural exposure to the same agent. The route of acquisition of the infection does not bear any relationship to the risk of secondary transmission. However, a suspected BW attack on any military deployed operation would have significant political, operational and medical implications. Medical factors will include the need for:
- a. Rapid and accurate identification of the agent in order to ensure appropriate prophylaxis of non-exposed personnel and treatment of those exposed.
- b. Provision of adequate medical resources in theatre. This will include diagnosis, treatment, specialist nursing, infection control and the ability, if required, to hold BW casualties.
 - c. Confirmation of the risk of secondary infection.
- 2. Aim. The aim of this paper is to identify the actions that need to be taken to prevent the spread of a contagious disease from a deployed theatre of operations to the home base from the aeromedical evacuation of casualties.
- 3. Scope. This paper is concerned only with medical issues. It does not address the need or provision for forensic sampling. Detailed guidance on Restriction of Movement (ROM) is addressed in Appendix 7.
- 4. Attack Indicators. The following are indicators that BW agents have been used:
 - a. Intelligence.
 - b. Activation of detectors.
- c. Medical surveillance. For example, the sudden onset of illness with large numbers of personnel military or civilians or animals, unusual distribution or types of illness in the theatre of operations (e.g., localized or widespread, multiple foci).
- 5. Restriction of Movement (ROM). In the event of a suspected BW incident, the theatre commander may impose ROM along with other disease-control measures. The presence of positive intelligence (i.e., both the possession of BW agents by a nation of concern and an indication of intention to use) together with the occurrence of 4b or 4c above will almost certainly be treated as a BW attack until proved otherwise. Where a contagious disease is

suspected it will, <u>at a minimum</u>, be necessary to ensure that those returning home from a theatre of operations (including casualties) are the subjects of rigorous follow-up health surveillance. As a general rule, the movement of personnel and casualties back to the home base should be minimized until any disease is definitively diagnosed. Where ROM is implemented, a decision to evacuate casualties from the theatre of operation will need to be taken in consultation with the theatre commander and operational HQ. Any decision to evacuate casualties needs to be balanced against the risk of spreading a contagious disease to the home base and the needs of the casualties. ROM may be applied to the movement of casualties.

- 6. Actions. A simple decision-making tool covering the steps that need to be taken in order to avoid the inadvertent spread a contagious disease back to the home base is attached at Addendum A, Evidence from BW Sensor Alarms (just ahead of this appendix). This tool will be implemented under ROM or where there is suspicion of a contagious disease affecting personnel in-theatre. Up to three steps may be required to accurately diagnose a disease:
- Step 1: Diagnosis in theatre. Ideally, disease diagnosis will be made in theatre. This might be based on clinical observations of symptomatic cases or the analysis of suitable samples from symptomatic patients using in-theatre analysis capabilities.
- Step 2: Returning Specimens to the UK/US. If diagnosis cannot be made within the theatre of operations, specimens must be sent out of theatre. Take samples only from patients who are symptomatic. There is no benefit in taking samples from healthy individuals who may have been exposed to a contagious agent. Consider sample collection from corpses. Veterinary and/or environmental-health assistance may be required to collect samples from dead animals.
 - Step 3: Evacuation of an Index Case to the UK/US.
- a. The initial clinical picture of natural and weaponized agents is often non-specific and the analysis of samples may not result in a definitive diagnosis. A patient may therefore need to be moved to a reference facility for confirmation of diagnosis and guidance on treatment. In order to facilitate the diagnosis process, it may be necessary to evacuate an index case along with the transmission of samples at Step 2.
- b. Evacuation of an Index Case–Mechanism. The patient should be transported in a patient Air Transportable Isolator (ATI) chamber because of:
 - (1) Risk of airborne transmission.
 - (2) Risk of contamination of the airframe.
- c. Selection of Index Patient. Senior medical officer in charge of the patient, following consultation with the theatre medical commander, will make the choice of individual. In general, the patient should be representative of the clinical syndrome involved, early in the natural history of the illness and physically accessible to the ATI team. Ideally the patient

should be self-caring and unlikely to deteriorate in the time it takes to reach the reference facility. The evacuation of an index case will comply, whenever possible, with IATA and WHO Regulations.

- Step 4: Risk Assessment. The implementation of the above steps should safeguard the home base against inadvertent spread of contagious disease from the aeromedical evacuation of casualties. Once an accurate diagnosis has been made, the theatre medical Commander and his staff will need to:
- a. Identify the health protection options and medical resources required, including the most appropriate prophylaxis and treatments for any personnel who may have been exposed and/or infected with a contagious disease. This process will assist the commander in deciding what force health protection measures, including the use of ROM, need be implemented in-theatre.
- b. Carry out an assessment of the risks of evacuating casualties, some of who may be contagious, back to the home base. It may be possible to evacuate casualties with appropriate precautions in place, however where it is judged that the risks of doing this are too great treatment will need to be delivered in theatre.
- 7. Personnel & Casualties Already Evacuated. It is possible that Steps 1-4 above will take several days to effect. By the time a definitive diagnosis is obtained, some casualties may already have been evacuated to the home base. Where a contagious disease is confirmed, evacuated casualties and contacts will need to be the subject of rigorous medical surveillance.

Annex J J-28 29 Sep 02

APPENDIX J-7

Advisory Committee on Immunization Practices (ACIP) Guidelines:

Part 1, Summary of October 2002 ACIP Smallpox Vaccination Recommendations.

Approved by ACIP on October 18, 2002

Now under consideration by CDC and DHHS

http://www.bt.cdc.gov/agent/smallpox/vaccination/acip-recs-oct2002.asp

Part 2, Draft Supplemental Recommendation of the ACIP, Use of Smallpox (Vaccinia) Vaccine, June 2002.

Draft approved by ACIP on June 20, 2002

Now under consideration by CDC and DHHS

http://www.bt.cdc.gov/agent/smallpox/vaccination/acip-guidelines.asp

Part 3, ACIP Recommendations for Use of Smallpox Vaccine, June 2001

Advisory Committee on Immunization Practices (ACIP).

Vaccinia (smallpox) vaccine.

MMWR—Morbidity & Mortality Weekly Report 2001;50(RR-10)(June 22):1-25.

http://www.cdc.gov/mmwr/PDF/rr/rr5010.pdf.

The documents appear on following pages.